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DESCRIPTION

NEGATIVE PHOTSENSITIVE RESIN COMPOSITION, NEGATIVE
PHOTSENSITIVE DRY FILM AND METHOD OF FORMING PATTERN

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TECHNICAL FIELD

The present invention relates to novel negative
photosensitive resin compositions, negative photosensitive
dry films and methods of forming patterns.

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BACKGROUND ART

Negative photosensitive resin compositions have
been widely used in paints, inks, adhesives, resist
materials, printing plate materials, information recording
materials, materials for producing relief images, etc.,
because of their excellent characteristics such as non-
polluting properties, resource- and energy-saving
properties, high production efficiency and the like.

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Negative photosensitive resin compositions are
used in various applications, by utilizing the solubility
difference occurring between exposed and unexposed parts.
Conventionally, exposure of these compositions is carried
out using visible light with an emission spectrum
wavelength of, for example, 488 nm or 532 nm. However,
conventional negative photosensitive resin compositions
are not sufficiently sensitive to visible light,

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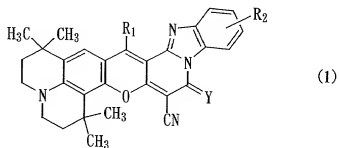
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(C) a photosensitizer which is a benzopyran condensed ring compound capable of increasing photosensitivity to visible light with a wavelength of 480 nm or more.

2. A composition according to item 1, wherein
5 the proportion of the photoacid generator (B) is about 0.01 to 10 parts by weight per 100 parts by weight of the resin (A).

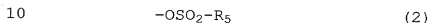
3. A composition according to item 1, wherein
10 the photosensitizer (C) is a benzopyran condensed ring compound represented by Formula (1)



wherein R_1 is hydrogen, halogen, cyano, trifluoromethyl,
15 carboxyl or carboxylic acid ester, R_2 is hydrogen, alkyl, alkoxy, cyano, trifluoromethyl, sulfoxy or halogen, and Y is NH or O.

4. A composition according to item 1, wherein
20 the proportion of the photosensitizer (C) is about 0.1 to 10 parts by weight per 100 parts by weight of the total amount of the resin (A) and photoacid generator (B).

5. A composition according to item 1 which further comprises, as an photoacid proliferating agent (D), an organic acid ester (a) and/or a crosslinked carbocyclic compound (b) containing a crosslinked carbocyclic skeleton which has a hydroxyl group or groups bonded to any of the crosslinked carbocyclic rings and, at a carbon atom or atoms adjacent to the hydroxyl-bearing carbon atom or atoms, a sulfonate group represented by Formula (2)



wherein R_5 is acyl, aliphatic hydrocarbon, polycyclic aromatic hydrocarbon or a heterocyclic group.

6. A composition according to item 1 which is an organic solvent-based resin composition.

15 7. A composition according to item 1 which is an aqueous resin composition.

8. A negative photosensitive dry film prepared by applying a negative photosensitive resin composition according to item 1 to a surface of support film, followed
20 by drying, to form a negative photosensitive resin layer.

9. A method of forming a pattern comprising the steps of:

(1) applying a negative photosensitive resin composition according to item 1 to a substrate, followed by drying, to
25 form a negative photosensitive resin coating,

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(2) irradiating the resin coating with visible light directly or through a mask so as to obtain a desired pattern, and

(3) removing the unirradiated part of the negative photosensitive resin coating by development to form a resist pattern coating.

10. A method of forming a pattern comprising the steps of:

(1') attaching a negative photosensitive dry film according to item 8 to a substrate so that the photosensitive resin layer of the dry film is in contact with the substrate to form a negative photosensitive resin coating, and optionally peeling off the support film of the dry film,

(2) irradiating the resin coating with visible light directly or through a mask so as to obtain a desired pattern, and

(3') peeling off the support film of the dry film when the support film has not been peeled off, and removing the unirradiated part of the negative photosensitive resin coating by development to form a resist pattern coating.

The present inventors carried out intensive research to overcome the drawbacks of the prior art, and found that the drawbacks can be overcome by using, as a photosensitizer, a benzopyran condensed ring compound

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capable of increasing photosensitivity to visible light with a wavelength of 480 nm or more. The present invention has been accomplished based on this finding.

5 Negative photosensitive resin composition

The negative resin composition of the present invention comprises (A) a photocurable resin having a photosensitive group or groups crosslinkable by light irradiation, (B) a photoacid generator and (C) a
10 photosensitizer which is a benzopyran condensed ring compound capable of increasing sensitivity to visible light with a wavelength of 480 nm or more.

The photocurable resin (A) having a photosensitive group or groups crosslinkable by light
15 irradiation for use in the invention is a compound which is cured and rendered insoluble by polymerization, etherification reaction, pinacol rearrangement, silanol dehydration reaction, intramolecular dehydration
condensation reaction, hydrolysis condensation reaction or
20 like reaction using, as a catalyst, an acid generated from the acid generator (B) by light irradiation.

Any conventional photocurable resin can be used as the photocurable resin (A) without limitation.

Resins usable as the resin (A) include, for
25 example, glycidyl ether epoxy compounds, alicyclic epoxy

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compounds, vinyl ether compounds, alkoxyallene compounds, oxetane compounds, ketene acetal compounds, bicyclo ortho ester compounds, lactone compounds, aromatic vinyl compounds, heterocyclic vinyl compounds, melamine

- 5 compounds and other aromatic compounds. These compounds may be low-molecular-weight or high-molecular-weight ones as long as they are curable by an acid. These compounds have a photosensitive group or groups such as epoxy, vinyl ether, allene, oxetane, acetal, ortho ester, lactone ring, vinyl, triazine ring and the like.

Glycidyl ether epoxy compounds include, for example, bisphenol A diglycidyl ether, (poly)ethylene glycol diglycidyl ether and trimethylolpropane diglycidyl ether.

- 15 Alicyclic epoxy compounds include, for example, those disclosed in J. Polym. Sci.: Part A: Polym. Chem. 28, 479 (1990). Specific examples include 3,4-epoxycyclohexylmethyl-3,4-epoxycyclohexane carboxylate, dicyclopentadiene dioxide, epoxycyclohexene carboxylic acid ethylene glycol diester, and 1,3-bis[2-(3(7-oxabicyclo[4.1.0]heptyl)ethyl)-tetramethyldisiloxane.

- 25 Vinyl ether compounds include, for example, those disclosed in J. Polym. Sci.: Part A: Polym. Chem. 34, 2051 (1996). Specific examples include butylene glycol divinyl ether, trimethylolpropane di(1-propenyl)

methylether, trimethylolpropane di(1-propenyl) butylether, trimethylolpropane di(1-propenyl) octylether, trimethylolpropane di(1-propenyl) phenylether, trimethylolpropane di(1-propenyl) ether acetate, 5 trimethylolpropane di(1-propenyl) ether acrylate and trimethylolpropane di(1-propenyl) N-butyl carbonate.

Alkoxy allene compounds include, for example, those disclosed in J. Polym. Sci.: Part A: Polym. Chem. 33, 2493 (1995). Specific examples include dodecyl allene (DA), 10 diethylene glycol diallene (DEGA), triethylene glycol diallene (TEGA), 1-tetrahydrofurfuryl allene ether (THFA), N-hexyloxy-1,2-propadiene (HA), 1,4-di-N-butoxy-1,2-butadiene (DBB), 1,4-diethoxy-1,2-butadiene, and N-hexylpropazyl ether (HPE).

15 Oxetane compounds include, for example, those disclosed in J. Polym. Sci.: Part A: Polym. Chem. 33, 1807 (1995). Specific examples include 3-ethyl-3-phenoxy-methyl-oxetane, phenoxy-methyloxetane, methoxy-methyloxetane and 3-methyl-3-methoxy-methyl-oxetane.

20 Ketene acetal compounds include, for example, those disclosed in J. Polym. Sci.: Part A: Polym. Chem. 34, 3091 (1996). Specific examples include 2-propylidene-4,5-dimethyl-1,3-dioxolane, 2-propylidene-4-methyl-1,3-dioxolane and 3,9-dimethylidene-2,4,8,10-

25 tetraoxapir[5.5]undecane.

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Bicyclo ortho ester compounds include, for example, those disclosed in J. Polym. Sci.: Polym. Lett. Ed. 23, 359 (1985). Specific examples include 1-phenyl-4-ethyl-2,6,7-trioxabicyclo[2.2.2]octane.

5 Lactone compounds include, for example, propiolactone, butyrolactone, γ -valerolactone, γ -caprolactone, γ -caprylolactone, γ -laurylolactone and coumarin.

10 Aromatic vinyl compounds include methoxy- α -methylstyrene and the like.

Heterocyclic vinyl compounds include vinyl carbazole and the like.

Melamine compounds include, for example, hexamethylolmelamine and hexamethoxymelamine.

15 Other aromatic compounds include, for example, copolymers of p-vinylphenol and p-vinylbenzyl acetate, trimethylolbenzene and tri(acetoxycarbonylmethyl)benzene.

20 The photoacid generator (B) for use in the negative photosensitive resin composition of the invention is a compound capable of generating an acid when exposed. The generated acid acts as a catalyst to cure the photocurable resin (A). Known photoacid generators are usable, which include sulfonium salts, ammonium salts, phosphonium salts, iodonium salts, selenium salts and like
25 onium salts; iron-allene complexes; silanol-metal chelate

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complexes; triazine compounds; diazide naphthoquinone compounds; sulfonic acid esters; sulfonic acid imide esters; and halogen compounds.

Among these compounds, preferred are onium salts, 5 sulfonic acid imide esters and the like. Also usable are photoacid generators disclosed in Japanese Unexamined Patent Publication No. 1995-146552 and Japanese Patent Application No. 1997-289218.

Specific examples of iodonium salts include 10 chlorides, bromides, borofluorides, hexafluorophosphate salts or hexafluoroarsenate salts of diphenyliodonium, ditolyliodonium, phenyl(p-anisyl)iodonium, bis(m-nitrophenyl)iodonium, bis(p-chlorophenyl)iodonium or like iodonium.

15 Specific examples of sulfonium salts include triarylsulfonium salts and dialkyl-4-hydroxysulfonium salts.

Specific examples of phosphonium salts include triarylphosphonium salts.

20 Specific examples of sulfonic acid esters include benzoin tosylate, pyrogallol trimesylate, o-nitrobenzyl tosylate, 2,5-dinitrobenzyl tosylate, N-tosyl phthalic acid imide, α -cyanobenzylidene tosyl amine and p-nitrobenzil-9,10-diethoxyanthracene-2-sulfonate.

25 Specific examples of sulfonic acid imide esters

include imide sulfonate.

Specific examples of silanol-metal chelate complexes include silanol-aluminium complexes.

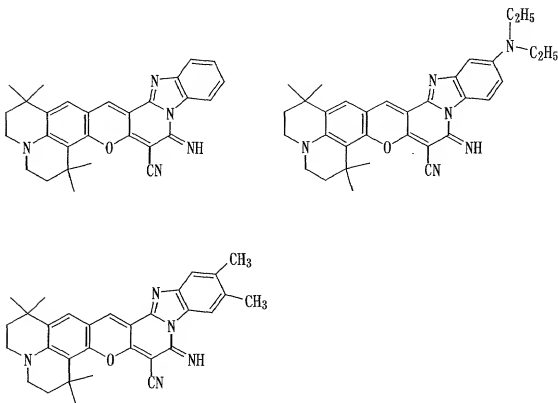
Commercial products of photoacid generators are,
5 for example, Cyracure UVI-6970, Cyracure UVI-6974,
Cyracure UVI-6990 and Cyracure UVI-6950 (tradenames of
Union Carbide Corp. (U.S.)), Irgacure 261 (a tradename of
Ciba Specialty Chemicals), SP-150 and SP-170 (tradenames
of Asahi Denka Kogyo K.K.), CG-24-61 (a tradename of Ciba
10 Specialty Chemicals), DAICAT-11 (a tradename of Daicel
Chemical Industries, Ltd.), CI-2734, CI-2758 and CI-2855
(all tradenames of Nippon Soda Co., Ltd.), PI-2074 (a
tradename of Rhone Poulenc SA, pentafluorophenylborate
tolyl cumyl iodonium salt), FFC509 (a tradename of 3M),
15 BBI102 and NAI-105 (tradenames of Midori Kagaku Co., Ltd.),
etc.

The photoacid generator (B) can be used as mixed
with the resin (A) or as bonded to the resin (A).

Resins containing the photoacid generator (B)
20 incorporated into the skeleton of the resin (A) are, for
example, resins which generate acid groups when exposed,
to thereby enable alkali development. Examples of such
resins include those in which a naphthoquinone diazide
sulfonic acid is bonded to an acrylic resin or like resin
25 containing ion-forming groups via a sulfonic acid ester

The alkyl group represented by R_2 include methyl, ethyl, butyl, octyl, dodecyl and octadecyl. Examples of the alkoxy group represented by R_2 include methoxy and ethoxy.

As specific examples of the benzopyran condensed 5 ring compound of Formula (1), the following compounds can be mentioned.



10 In the present invention, photosensitizing dyes can be used as required, in addition to the photosensitizer (C). Examples of photosensitizing dyes include thioxanthene dyes, xanthene dyes, ketone dyes,

thiopyrylium salt dyes, base styryl dyes, merocyanine dyes, 3-substituted coumarin dyes, 3,4-substituted coumarin dyes, cyanine dyes, acridine dyes, thiazine dyes, phenothiazine dyes, anthracene dyes, coronene dyes, benzanthracene dye, 5 perylene dyes, merocyanine dyes, ketocoumarin dyes, fumarine dyes, borate dyes and other photosensitizing dyes. Usable borate photosensitizing dyes include, for example, those shown in Japanese Unexamined Patent Publications No. 1993-241338, No. 1995-5685 and No. 1995-225474.

10 The proportion of the photosensitizer (C) for use in the composition of the invention is preferably about 0.1 to 10 parts by weight, especially about 0.3 to 5 parts by weight, per 100 parts by weight of the total amount of the resin (A) and photoacid generator (B).

15 The composition of the invention may further comprise a photoacid proliferating agent (D), if necessary.

Preferred as the photoacid proliferating agent (D) are an organic acid ester (a) and a crosslinked carbocyclic compound (d) containing a crosslinked 20 carbocyclic skeleton which has a hydroxyl group or groups bonded to any of the crosslinked carbocyclic rings and, on a carbon atom or atoms adjacent to the hydroxyl-bearing carbon atom or atoms, a sulfonate group represented by Formula (2)

25



(2)

wherein R_5 is acyl, aliphatic hydrocarbon, polycyclic aromatic hydrocarbon or a heterocyclic group. The aliphatic hydrocarbon group represented by R_5 in Formula (2) may be, for example, chain or cyclic (including crosslinked carbocyclic) alkyl or alkenyl. The aliphatic hydrocarbon group has usually 1 to 12, preferably 1 to 8, carbon atoms. The polycyclic aromatic hydrocarbon group may be, for example, naphthyl, biphenyl or naphthylmethyl. The heterocyclic group may have a monocyclic or polycyclic structure, and may be any heterocyclic group derived from various known heterocyclic compounds. These aliphatic hydrocarbon group, polycyclic aromatic hydrocarbon group and heterocyclic group may be substituted by halogen, hydrocarbonoxy, amino, substituted amino or the like.

Examples of acyl groups include formyl, acetyl, benzoyl and cinnamoyl. Examples of aliphatic hydrocarbon groups and polycyclic aromatic hydrocarbon groups include methyl, ethyl, propyl, butyl, hexyl and like alkyl groups; vinyl, propenyl, allyl and like alkenyl groups;

cyclohexyl, cyclooctyl, bicyclo hydrocarbon, tricyclo hydrocarbon and like cycloalkyl groups; naphthyl, naphthylmethyl, biphenyl and like aryl or arylalkyl groups; and these groups in a substituted form. Specific examples of heterocyclic groups include those derived from various heterocyclic compounds, such as furan, thiophene,

pyrrole, benzofuran, thionaphthene, indole, carbazole and like five-membered ring compounds containing one hetero atom, and condensed ring compounds thereof; oxazole, thiazole, pyrazole and like five-membered compounds having
5 two hetero atoms, and condensed ring compounds thereof; pyran, pyrone, coumarine, pyridine, quinoline, isoquinoline, acridine and like six-membered compound having one hetero atom, and condensed ring compounds thereof; and pyridazine, pyrimidine, pyrazine, phthalazine
10 and like six-membered compounds containing two hetero atoms, and condensed ring compounds thereof.

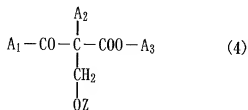
The photoacid proliferating agent (D), when used in combination with the photoacid generator (C), is decomposed by an acid released from the photoacid
15 generator (C) by light irradiation to thereby produce a free acid. The produced free acid further decomposes the photoacid proliferating agent so that a free acid is further produced. Thus, the photoacid proliferating agent is decomposed in a chain-like manner by light irradiation,
20 thereby forming numerous free acid molecules.

The organic acid ester (a) is a compound substituted by a residue of a relatively strong acid and capable of readily producing an acid by elimination in the presence of a photoacid generator. Specifically stated,
25 the ester (a) is decomposed by an acid generated from a

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photoacid generator and produces an acid (hereinafter referred to as "ZOH"). Since at least one acid molecule is produced by one reaction, the acid concentration is acceleratedly increased as the reactions proceed, and thus the composition is remarkably improved in photosensitivity. Preferred acid strength of the generated acid (ZOH) is an acid dissociation constant (pKa) not greater than 3, especially not greater than 2. A weaker acid is incapable of causing elimination reaction by an acid catalyst. As examples of the acid (ZOH), there can be mentioned dichloroacetic acid, trichloroacetic acid, methanesulfonic acid, ethanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, naphthalenesulfonic acid, phenylphosphonic acid and the like.

The first example of the photoacid proliferating agent (D) is an organic acid ester compound represented by Formula (4):

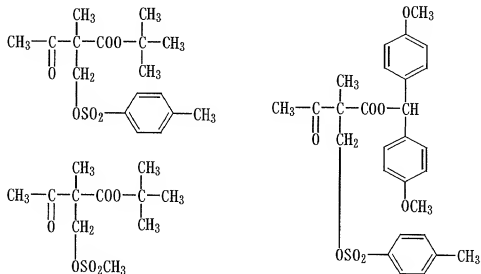


wherein A₁ is C₁ to C₆ alkyl or aryl, A₂ is C₁ to C₆ alkyl, A₃ is bis(p-alkoxyphenyl)methyl, 2-alkyl-2-propyl, 2-aryl-2-propyl, cyclohexyl or tetrahydropyranyl, Z is a residue

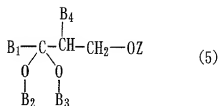
of an acid represented by the formula ZOH and having a pKa not greater than 3.

In Formula (4), the C₁ to C₆ alkyl groups represented by A₁ and A₂ each may be, for example, methyl, ethyl, propyl or the like. The aryl group represented by A₁ may be, for example, phenyl or the like.

When an acid acts on the compound of Formula (4), the ester group decomposes into carboxylic acid. Further, decarboxylation is caused to eliminate the acid (ZOH). Specific examples of the compound of Formula (4) are as follows.



The second example of the photoacid proliferating agent (D) is an acetal- or ketal-containing organic acid ester represented by Formula (5):

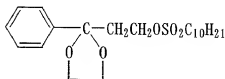
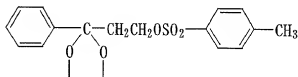


wherein Z is a residue of an acid represented by ZOH and having a pKa not greater than 3, B₁ is hydrogen, alkyl or aryl, B₂ and B₃ are each methyl or ethyl, or B₂ and B₃ are bonded together to form ethylene or propylene, and B₄ is hydrogen or methyl.

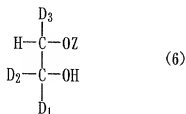
The alkyl group represented by B₁ in Formula (5) may be, for example, methyl, ethyl, propyl or the like.

The aryl group represented by B₁ may be, for example, phenyl or the like.

In the compound of Formula (5), acetal or ketal is decomposed by an acid into β-aldehyde or ketone, from which ZOH is easily eliminated. Specific examples of the compound of Formula (5) are as follows:



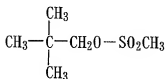
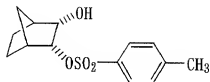
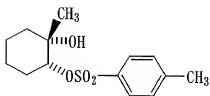
The third example of the photoacid proliferating agent (D) is an organic acid ester represented by Formula (6):



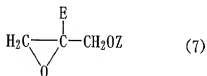
5 wherein Z is a residue of an acid represented by ZOH and having a pKa not greater than 3, D₁ and D₃ are each C₁ to C₆ alkyl, or D₁ and D₃ are bonded together to form
10 alkylene or substituted alkylene constituting an alicyclic structure, and D₂ is hydrogen, C₁ to C₆ alkyl or aryl.

In Formula (6), the C₁ to C₆ alkyl groups represented by D₁, D₂ and D₃ each may be methyl, ethyl, propyl or the like. The alkylene or substituted alkylene
15 group formed by D₁ and D₃ to constitute an alicyclic structure may be, for example, heptylene or the like. An example of the aryl group represented by D₂ is phenyl or the like.

It is presumed that, in the compound of Formula (6), the hydroxyl group is eliminated by an acid catalyst
20 to form a carbocation, undergoes hydrogen transfer and then generates ZOH. Specific examples of the compound of Formula (6) are as follows.



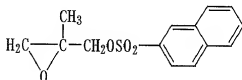
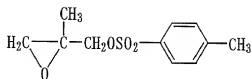
- The fourth example of the photoacid
proliferating agent (D) is an epoxy ring-containing
5 organic acid ester represented by Formula (7).



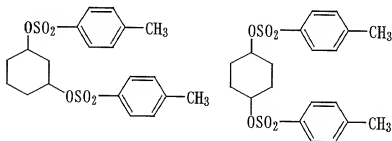
- wherein Z is a residue of an acid represented by ZOH and
having a pKa not greater than 3, and E is C₁ to C₆ alkyl
10 or phenyl.

Examples of the C₁ to C₆ alkyl group represented
by E in Formula (7) include methyl, ethyl and propyl.

- It is presumed that, when an acid acts on the
compound of Formula (7), the epoxy ring opens to form a
15 cation at the β-carbon and an organic acid is generated as
a result of hydrogen transfer. Specific examples of the
compound of Formula (7) include the following:



Also usable as the organic acid ester are, for example, cis-3-(p-toluenesulfonyloxy)-2-pinanol as a monofunctional compound, and the following compounds as multifunctional compounds.



The organic acid ester is stable at room temperature as long as no acid acts thereon. A specific acid strength is required for decomposition of the compound in the presence of an acid catalyst. A preferred acid strength is a pKa not greater than 3, especially not greater than 2. A weak acid having a pKa over 3 is incapable of causing reaction of the photoacid

proliferating agent.

The crosslinked carbocyclic compound (b) for use as the photoacid proliferating agent (D) contains a crosslinked carbocyclic skeleton which has a hydroxyl group or groups bonded to any of the crosslinked carbon rings and, on a carbon atom or atoms adjacent to the hydroxyl-bearing carbon atom or atoms, a sulfonate group represented by Formula (2).

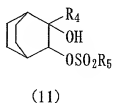
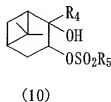
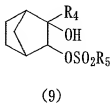
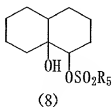
The compound (b) is decomposed by acid catalytic reaction and generates an acid (R_5SO_3H). One acid molecule is generated by one reaction, so that acid molecules are increased acceleratedly as the reactions proceed. For inducing self-decomposition of the generated acid, the acid strength of the generated acid is an acid dissociation constant (pK_a) not greater than 3, especially not greater than 2. A weaker acid is not incapable of inducing self-decomposition. Examples of acids (R_5SO_3H) released by such reactions include methanesulfonic acid, ethanesulfonic acid, propanesulfonic acid, butanesulfonic acid, pentanesulfonic acid, hexanesulfonic acid, heptanesulfonic acid, octanesulfonic acid, cyclohexanesulfonic acid, camphorsulfonic acid, trifluoromethanesulfonic acid, 2,2,2-trifluoroethanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, p-bromobenzenesulfonic acid, p-

nitrobenzenesulfonic acid, 2-thiophenesulfonic acid, 1-naphthalenesulfonic acid and 2-naphthalenesulfonic acid.

The compound (b) to be used as a photoacid proliferating agent can be decomposed in a chain-like manner by heating in the presence of an acid generated from a photoacid generator.

The compound (b) is characterized by having a crosslinked carbocyclic skeleton structure, such as a crosslinked structure comprising more than one, usually 2 to 6, preferably 2 to 3, carbocyclic rings. The crosslinked carbocyclic rings may have substituent(s) such as methyl, ethyl, propyl or like C_1 to C_6 , preferably C_1 to C_3 lower alkyl, or unsaturated bond(s) such as double bond. The crosslinked carbocyclic rings have, within the molecule, crosslink(s) which rigidify the molecule, thus giving a photoacid proliferating agent with improved thermal stability.

As preferred examples of the compound (b), compounds represented by the following Formulas (8) to (11) can be mentioned.



- In the above Formulas, R_5 is as defined above, and R_4 is hydrogen, aliphatic hydrocarbon or aromatic hydrocarbon. Aliphatic hydrocarbon groups include chain or cyclic (including crosslinked cyclic) alkyl or alkenyl.
- 5 Aromatic hydrocarbon groups include aryl, arylalkyl or the like. The aliphatic hydrocarbon and aromatic hydrocarbon groups may be substituted by halogen, alkoxy, amino, substituted amino or the like.

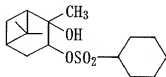
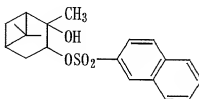
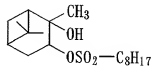
- The aliphatic hydrocarbon group has 1 to 12, preferably 1 to 8, carbon atoms. The aromatic hydrocarbon group may have a monocyclic or polycyclic structure. R_4 is preferably aliphatic hydrocarbon or aromatic hydrocarbon.
- 10

- Examples of alkyl groups include methyl, ethyl and propyl. Examples of alkenyl groups include propenyl. Examples of aryl groups include phenyl. The halogen atom may be, for example, fluorine or chlorine.
- 15

- The bicyclo compound of Formula (8) (a decalin derivative) has crosslinks at the 1- and 6- positions.
- 20 The bicyclo compounds of Formulas (9) and (11) have crosslinks at the 1- and 4-positions. The bicyclo compound of Formula (10) has crosslinks at the 1- and 3- positions. Therefore, in these bicyclo compounds, conformational change of the cyclohexane ring is highly suppressed and thus the ring structure shows rigidity.
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Further, the composition of the invention may optionally contain an adhesion promoter; hydroquinone, 2,6-di-t-butyl-p-cresol, N,N-diphenyl-p-phenylenediamine or like polymerization inhibitor; a rubber, a vinyl polymer, an unsaturated group-containing vinyl polymer or like particulate organic resin; a coloring pigment, an extender pigment and like pigment; a dye; cobalt oxide or

5 The adhesion promoter is used to improve
adhesion of a coating to a substrate. Specific examples
of adhesion promoters include tetrazole, 1-phenyltetrazole,
5-aminotetrazole, 5-amino-1-methyltetrazole, 5-amino-2-
phenyltetrazole, 5-mercapto-1-phenyltetrazole, 5-mercapto-
0 1-methyltetrazole, 5-methylthiotetrazole, 5-chloro-1-
phenyl-1H-tetrazole and like tetrazoles.

The negative photosensitive resin composition of the invention can be prepared by mixing the ingredients in a routine manner. Use of a solvent is optional in the preparation of the composition.

The organic solvent-based negative photosensitive resin composition of the invention can be prepared by dissolving or dispersing the ingredients in an organic solvent such as a ketone, an ester, an ether, a cellosolve, an aromatic hydrocarbon, an alcohol or a halogenated hydrocarbon, in a routine manner. The organic solvent-based composition has a solid concentration of preferably about 1 to 50 wt.%.

The aqueous negative photosensitive resin composition of the invention can be prepared by dissolving

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Negative photosensitive dry film

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the support film by roller coating, roll coater coating, spray coating, gravure coating or like coating process, and then drying the applied composition to volatilize water and organic solvent. During drying, the applied composition may be heated, if necessary.

The support film is usually about 5 to 100 μm thick, and the negative photosensitive resin layer is usually about 1 to 100 μm thick.

Usually, the negative photosensitive dry film is attached to a surface of a substrate and irradiated with light before or after peeling off the support film.

Use of the negative photosensitive resin composition and negative photosensitive dry film

The negative photosensitive resin composition and negative photosensitive dry film of the invention are excellent in sensitivity to visible light, storage stability and other properties. Thus, the composition and dry film are widely useful in conventional applications of photosensitive materials, such as paints, inks, adhesives, resist materials, printing plate materials, information recording materials, materials for producing relief images, etc.

Resist materials include, for example, photoresists, solder resists and plating resists.

d)

- glass-epoxy plates, polyethylene terephthalate films, polyimide films and like plastic films and plastic plates; these plastic plates and plastic films having a conductive layer formed by attaching foil of copper, aluminium or
- 5 like metal; substrates having a conductive layer formed by vacuum deposition, chemical vapor deposition, plating or like process using copper, nickel, silver or like metal or a compound of a conductive oxide (typically indium tin oxide (ITO)) or the like; plastic plates and plastic films
- 10 provided with a through hole portion, whose surface and through hole portion are covered with a conductive layer; and metal plates such as copper plates.

- The resin composition can be applied to a substrate by a process such as roller coating, roll coater
- 15 coating, spin coater coating, curtain roll coater coating, spray coating, electrostatic coating, dip coating, silk printing or spin coating.

- After applying the organic solvent-based or aqueous negative photosensitive resin composition to a
- 20 substrate, the composition is optionally set and dried at about 50 to 130°C to form a negative photosensitive resin coating.

- The photosensitive resin coating thus formed preferably has a dry thickness of about 0.5 to 100 μm ,
- 25 especially about 1 to 50 μm .

The photosensitive resin coating may be provided with a cover coat before curing by visible light exposure. The cover coat layer blocks oxygen in air and protects radicals generated by exposure from being inactivated by oxygen, so that the photosensitive resin coating is cured smoothly when exposed.

The cover coat layer can be formed by, for example, covering the surface of the photosensitive resin coating with a resin film made of a polyester resin (e.g., polyethylene terephthalate), an acrylic resin, polyethylene, a polyvinyl chloride resin (film thickness: about 1 to 70 μm).

Alternatively, the cover coat layer can be formed by applying an aqueous coating liquid obtained by dissolving or dispersing, in water, polyvinyl alcohol; partially saponified polyvinyl acetate; a polyvinyl alcohol-vinyl acetate copolymer; a partially saponified polyvinyl acetate-vinyl acetate copolymer; polyvinylpyrrolidone; pullulan and like water-soluble polysaccharide polymer; or an acrylic resin, a polyester resin, a vinyl resin, an epoxy resin or like aqueous resin containing basic group(s), acid group(s) or base(s), to the surface of the photosensitive resin coating to a dry thickness of about 0.5 to 5 μm , followed by drying. The cover coat layer is preferably removed after exposure of

5 disperses the resin in the aqueous coating liquid.

10 coating so that a desired resist pattern coating can be
formed in the subsequent step (3).

15 such as extra-high-pressure, high-pressure, medium-
pressure or low-pressure mercury lamps, chemical lamps,
carbon arc lamps, xenon lamps, metal halide lamps,
fluorescent lights, tungsten lamps and sunlight. Also
usable are various lasers that produce an oscillating beam
20 in the visible region. As laser sources, an argon laser
(oscillating beam at 488 nm) or SHG-YAG laser (532 nm)
are preferable since they have a stable high output.

25 is removed by development to form a resist pattern coating.

The development can be preferably carried out by a liquid development process. In the liquid development process, the resin coating is, for example, sprayed with or dipped in a developer at about 10 to 80°C, preferably
5 at about 15 to 50°C, for about 1 to 60 minutes, preferably about 2 to 30 minutes, so as to form a pattern from the resin coating.

The liquid development process can be carried out, for example, using an alkaline developer when the
10 coating-forming resin contains acid group(s); or using an acid developer when the resin contains basic group(s); or using a water developer when the resin contains hydrophilic group(s); or using an organic solvent developer when the resin is soluble or dispersible in
15 organic solvents.

Alkaline developers include, for example, aqueous solutions of monomethylamine, dimethylamine, trimethylamine, monoethylamine, diethylamine, triethylamine, monoisopropylamine, diisopropylamine,
20 triisopropylamine, monobutylamine, dibutylamine, monoethanolamine, diethanolamine, triethanolamine, dimethylaminoethanol, diethylaminoethanol, ammonia, caustic soda, caustic potash, sodium metasilicate, potassium metasilicate, sodium carbonate,
25 tetraethylammonium hydroxide and the like.

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Acid developers include, for example, aqueous solutions of formic acid, crotonic acid, acetic acid, propionic acid, lactic acid, hydrochloric acid, sulfuric acid, nitric acid, phosphoric acid and the like.

- 5 It is usually preferable that these developers have an acid or alkali concentration of about 0.05 to 10 wt.%.

- Examples of organic solvents include hexane, heptane, octane, toluene, xylene, dichloromethane, 10 chloroform, carbon tetrachloride, trichloroethylene and like hydrocarbon solvents; methanol, ethanol, propanol, butanol and like alcohol solvents; diethyl ether, dipropyl ether, dibutyl ether, ethyl vinyl ether, dioxane, propylene oxide, tetrahydrofuran, cellosolve, methyl 15 cellosolve, butyl cellosolve, methyl carbitol, diethylene glycol monoethyl ether and like ether solvents; acetone, methyl ethyl ketone, methyl isobutyl ketone, isophorone, cyclohexanone and like ketone solvents; methyl acetate, ethyl acetate, propyl acetate, butyl acetate and like 20 ester solvents; pyridine, formamide, N,N-dimethyl formamide and like other solvents.

Thus, steps (1), (2) and (3) produce a desired resist pattern coating.

- The pattern forming method of the invention 25 which employs a negative photosensitive dry film comprises

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the steps of:

(1') attaching a negative photosensitive dry film of the invention to a substrate in such a manner that the photosensitive resin layer of the dry film is in contact with the substrate, to form a negative photosensitive resin coating, and optionally peeling off the support film of the dry film,

(2) irradiating the resin coating with visible light directly or through a mask so as to obtain a desired

pattern, and

(3') peeling off the support film of the dry film when the support film has not been peeled off, and removing the unirradiated part of the negative photosensitive resin coating by development to form a resist pattern coating.

In step (1'), a negative photosensitive dry film is attached to a substrate so that the photosensitive resin layer on the support film is in contact with the substrate, to form a negative photosensitive resin coating.

Useful substrates are as mentioned above. The support film may, but need not, be peeled off in this step. Drying is not necessary because a dry film is utilized.

Step (2) is as described above.

Step (3') is the same as step (3) except that the support film of the dry film is peeled off when the support film has not been peeled off in step (1').

Thus, steps (1'), (2) and (3') produce a desired resist pattern coating.

The method of the invention is capable of forming an extremely fine pattern with excellent contrast.

- 5 The method of the invention may be modified to be adapted to various applications, as long as it comprises the above steps.

- 10 The method finds applications in various industries including: electrical industries such as electrical components, lighting, electronic devices, semiconductors, printed circuits, electronic communication and electric power; physical industries such as measurement, optics, display, sound, control, automatic selling, signals and information recording; chemical, 15 metallurgical and fiber industries such as inorganic chemistry, organic chemistry, macromolecular chemistry, metallurgy and fibers; processing and transportation industries such as separation and mixing, metal working, plastic working, printing, containers and packaging; 20 daily necessities industries such as agricultural and marine products, foods, fermentation, household goods, health and recreation; and mechanical engineering industries.

- 25 Specific examples of applications in electrical industries include formation of black matrix insulating

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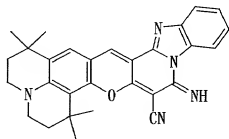
5 coatings, for color filters, fluorescent materials for display panels, hologram patterns, CD mastering, coils, etc. Specific examples of applications in physical industries include optical fiber processing, floppy disks, magnetic tapes, magnetic cards, optical components, wave absorbers, etc. Specific examples of applications in chemical, metallurgical and fiber industries include glass, cement, ceramics and like inorganic insulating materials. Specific examples of applications in processing and transportation industries include printed matters, original printing plates, diffraction grating, marking, barcodes, masks, filters, etching, defrosters, cement processing, stone processing, fiber processing, plastic processing, labels, etc. Specific examples of applications in daily necessities industries include carriers, cosmetic products, fermentation industries, etc. Specific examples of applications in mechanical engineering industries include micromachine components, etc.

BEST MODE FOR CARRYING OUT THE INVENTION

The following Examples are provided to illustrate the present invention in further detail. In these examples, parts and percentages are all by weight.

5 Example 1

An organic solvent-based negative photosensitive resin composition (I) was prepared by mixing 167 parts of a butylated melamine resin having a molecular weight of about 8,000 (a 60% solution of xylol/butanol (weight ratio: 50/50)), 1 part of Cyracure UVI-6990 (a tradename of Union Carbide Corporation (U.S.), a photoacid generator), 1 part of cis-3-(p-toluenesulfonyloxy)-2-pinanol and 1 part of the following benzopyran condensed ring compound.



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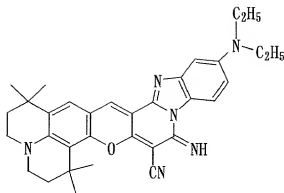
The composition (I) was applied to an aluminum plate coated with an epoxy primer, using a roll coater to a dry thickness of 20 μm and dried to form a negative photosensitive resin coating. The resin coating was irradiated with 300 mJ/cm^2 of visible light with a wavelength of 480 nm or more emitted from a metal halide

lamp covered with a filter, and heated at 110°C for 15 minutes for curing. The cured coating had a pencil hardness of H and high solvent resistance.

- Separately, a negative photosensitive resin
- 5 coating obtained in the same manner as above is irradiated with the same amount of the same visible light as above through a pattern mask, and heated at 110°C for 15 minutes for curing the irradiated part of the coating.
- Subsequently, the unirradiated part of the coating was
- 10 removed by development with methanol, thereby giving an excellent pattern coating.

Example 2

- An organic solvent-based negative photosensitive resin composition (II) was prepared by mixing 100 parts of
- 15 Nikalac MS-25 (a tradename of Sanwa Chemical Co., Ltd., a melamine resin having a nonvolatile content of 70%), 2 parts of CI-2855 (a tradename of Nippon Soda Co., Ltd., a photoacid generator), 1 part of cis-1-phenyl-2-p-toluenesulfonyloxy-1-cyclohexanol, 1 part of the following
- 20 benzopyran condensed ring compound and 20 parts of titanium dioxide.



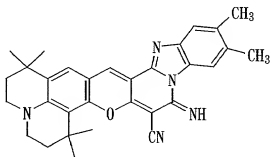
The composition (II) was applied to a surface-treated steel plate using a roll coater to a dry thickness of 10 μm and dried to form a negative photosensitive resin coating. The resin coating was irradiated with 400 mJ/cm^2 of visible light with a wavelength of 480 nm or more emitted from a metal halide lamp covered with a filter, and heated at 120°C for 10 minutes for curing. The cured coating had a pencil hardness of F.

Separately, a negative photosensitive resin coating obtained in the same manner as above was irradiated with the same amount of the same visible light as above through a pattern mask, and heated at 120°C for 10 minutes for curing the irradiated part of the coating. Subsequently, the unirradiated part of the coating was removed by development with methanol, giving an excellent pattern coating.

Example 3

An organic solvent-based negative photosensitive

resin composition (III) was prepared by mixing 140 parts of Phthalkyd 235-50 (a tradename of Hitachi Chemical Co., Ltd., an alkyd resin having a nonvolatile content of 50% and an oil length of 35), 55 parts of a butylated melamine resin having a molecular weight of about 5,000, 1 part of DAICAT-11 (a tradename of Daicel Chemical Industries, Ltd., a photoacid generator), 1 part of 1,4-di-p-toluenesulfonyloxy cyclohexane, 1 part of the following benzopyran condensed ring compound and 10 parts of calcium carbonate.

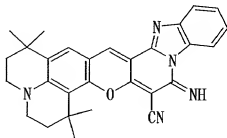


The composition (III) was applied to a primer-treated aluminium plate to a dry thickness of 30 μm by spray coating and dried to form a negative photosensitive resin coating. The resin coating was irradiated with 600 mJ/cm^2 of visible light with a wavelength of 480 nm or more emitted from a metal halide lamp covered with a filter, and heated at 110°C for 20 minutes for curing. The cured coating had a pencil strength of H.

Separately, a negative photosensitive resin coating prepared in the same manner as above was irradiated with the same amount of the same visible light through a pattern mask, and heated at 110°C for 20 minutes for curing the irradiated part of the coating. Subsequently, the unirradiated part of the coating was removed by development with methanol, giving an excellent pattern coating.

Example 4

An organic solvent-based negative photosensitive resin composition (IV) was prepared by mixing 117 parts of an acrylic resin vanish (nonvolatile content: 60%, a styrene/n-butyl methacrylate/n-butyl acrylate/hydroxybutyl acrylate copolymer having a hydroxyl value of 100 mg KOH/g and a molecular weight of 8,000), 55 parts of a melamine resin vanish (nonvolatile content: 55%, molecular weight: 8,000), 1.5 parts of DAICAT-11 (a tradename of Daicel Chemical Industries, Ltd., a photoacid generator), 1.5 part of 1,3-di-p-toluenesulfonyloxy cyclohexane and 1 part of the following condensed benzopyran compound.



The composition (IV) was applied to a glass plate using a roll coater to a dry thickness of 5 μm and dried to form a negative photosensitive resin coating.

The resin coating was irradiated with 200 mJ/cm^2 of visible light with a wavelength of 480 nm or more emitted from a metal halide lamp covered with a filter, and heated at 100°C for 15 minutes for curing. The cured coating had a pencil strength of H.

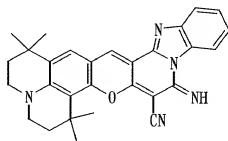
Separately, a negative photosensitive resin coating obtained in the same manner as above was irradiated with the same amount of the same visible light as above, and heated at 100°C for 15 minutes for curing the irradiated part of the coating. Subsequently, the unirradiated part of the coating was removed by development with methanol, giving an excellent pattern coating.

Example 5

An organic solvent-based negative photosensitive resin composition (V) was prepared by mixing 127 parts of an acrylic resin vanish (nonvolatile content: 55%, a styrene/methyl methacrylate/n-butyl acrylate/hydroxyethyl acrylate/acrylic acid copolymer having a hydroxyl value of 85 mgKOH/g , an acid value of 40 $\text{mg KOH}/\text{g}$ and a molecular weight of 10,000), 86 parts of Nikalac MS-35 (a tradename of Sanwa Chemical Co., Ltd., a melamine resin having a

nonvolatile content of 35%), 0.5 part of CI-2855 (a tradename of Nippon Soda Co., Ltd., a photoacid generator), 1 part of 1,4-di-p-toluenesulfonyloxy cyclohexane and 1 part of the following benzopyran condensed ring compound.

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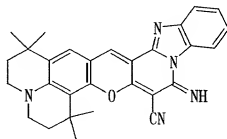
The composition (V) is applied to a polycarbonate plate by spray coating to a dry thickness of 20 μm and dried, giving a negative photosensitive resin coating. The resin coating was irradiated with 400 mJ/cm^2 of visible light with a wavelength of 480 nm or more emitted from a metal halide lamp covered with a filter, and heated at 100°C for 15 minutes for curing. The cured coating had a pencil hardness of H.

Separately, a negative photosensitive resin coating prepared in the same manner as above was irradiated with the same amount of the same visible light as above through a pattern mask, and heated at 100°C for 15 minutes for curing the irradiated part of the coating. Subsequently, the unirradiated part of the coating was

removed by development with a 1.5% aqueous sodium carbonate solution, giving an excellent pattern coating.

Example 6

5 An organic solvent-based negative photosensitive resin composition (VI) was prepared by mixing 150 parts of a water-soluble acrylic resin (emulsion type, nonvolatile content: 50%, hydroxyl value: 60 mg KOH/g, acid value: 45 mg KOH/g, molecular weight: 20,000), 25 parts of Cymel 303 (a tradename of Mitsui Cytec, Ltd., a water-soluble
10 melamine resin), 1 part of CI-2758 (a tradename, Nippon Soda Co., Ltd., a photoacid generator), 0.5 part of 1,4-di-p-toluenesulfonyloxy cyclohexane and 1 part of the following benzopyran condensed ring compound.



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The composition (VI) was applied to a surface-treated steel plate using a roll coater to a dry thickness of 20 μm and dried, giving a negative photosensitive resin coating. The resin coating was irradiated with 600 mJ/cm^2
20 of visible light with a wavelength of 480 nm or more emitted from a metal halide lamp covered with a filter,

Separately, a negative photosensitive resin coating prepared in the same manner as above was irradiated with the same amount of the same visible light as above through a pattern mask, and heated at 120°C for 15 minutes for curing the irradiated part of the coating. Subsequently, the unirradiated part of the coating was removed by development with a 1.5% aqueous sodium carbonate solution, giving an excellent pattern coating.